

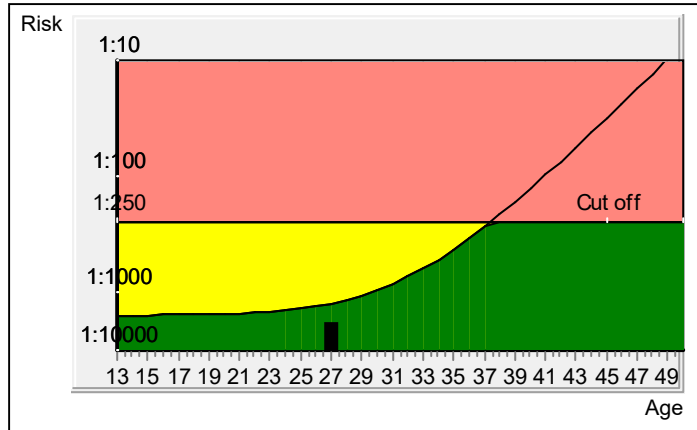
PRISCA 5.1.0.17

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Results for:
Mrs. PRIYANKA KUMARISample no
A0360273Date of report:
24-04-2024

Referring Doctors

Summary



Patient data

Age at delivery	26.9
WOP	14 + 3
Weight	45 kg
Patient ID	0482404230159
Ethnic origin	Asian

Risks at term

Biochemical risk for Tr.21	1:3675
Age risk:	1:1253
Neural tube defects risk	1:8236

For Mrs. PRIYANKA KUMARI, born on 08-11-1997, a screening test was performed on the 23-04-2024. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

TRISOMY 21 SCREENING

The calculated risk for Trisomy 21 is below the cut off which represents a low risk.

After the result of the Trisomy 21 test it is expected that among 3675 women with the same data, there is one woman with a trisomy 21 pregnancy and 3674 women with not affected pregnancies.

The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.

Please note that risk calculations are statistical approaches and have no diagnostic value!

MEASURED SERUM VALUES

	Value	Corr. MoMs
AFP	21.8 ng/mL	0.67
HCG	39787.8 mIU/mL	0.80
uE3	0.65 ng/mL	2.05

Gestation age 14+ 3

Method Scan

The MoMs have been corrected according to:
maternal weight
ethnic origin

TRISOMY 18 SCREENING

The calculated risk for trisomy 18 is < 1:10000, which indicates a low risk.

NEURAL TUBE DEFECTS (NTD) SCREENING

The corrected MoM AFP (0.67) is located in the low risk area for neural tube defects.

Risk above
Cut off

Risk above
Age risk

Risk below
Age risk